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This listing of the claims will replace all prior versions and listings of claims in the application:

Listing of the claims:

Claim 1 (original): A method for measuring enzymatically active Lipoprotein-associated Phospholipase A2 (Lp-PLA2) in a sample comprising:

- (a) contacting an immobilized binder, which specifically binds Lp-PLA2, with the sample;
- (b) washing the immobilized binder to remove an enzymatically active unbound material or an interfering substance(s);
- (c) contacting the bound Lp-PLA2 with a substrate converted to a detectable product in the presence of Lp-PLA2; and
- (d) measuring detectable product indicative of enzymatically active Lp-PLA2 in the sample.

Claim 2 (currently amended): The method of claim 1, wherein the sample is a serum sample, a plasma sample, or an EDTA treated plasma sample or an EDTA treated serum sample.

Claim 3 (original): The method of claim 1, wherein the immobilized binder is an antibody.

Claim 4 (original): The method of claim 3, wherein the antibody is a monoclonal antibody, a phage display antibody, or a polyclonal antibody.

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Claim 5-7 (canceled)

Claim 8 (original): The method of claim 1, wherein the substrate is selected from the group consisting of

$$\begin{array}{c|c} & & & & \\ & &$$

wherein,

X is selected from the group consisting of O, S, and -O(CO)-; R is selected from the group consisting of $(CH_2)_4CH_3$, $(CH_2)_6CH_3$, $(CH_2)_8CH_3$, $(CH_2)_{10}CH_3$, $(CH_2)_{12}CH_3$, $(CH_2)_{14}CH_3$, and $(CH_2)_7CH=CH(CH_2)_2CH_3$;

 Y_1 is selected from the group consisting of $(CO)_{1-2}$ and $(CH_2)_{2-7}$; and

 Y_2 is selected from the group consisting of CO and CH_2 ;

$$\begin{array}{c|c} & & & & & & \\ & & & & & \\ & & & & \\ & & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ &$$

wherein,

X is selected from the group consisting of O, S, and -O(CO)-; R is selected from the group consisting of $(CH_2)_4CH_3$, $(CH_2)_6CH_3$, $(CH_2)_8CH_3$, $(CH_2)_{10}CH_3$, $(CH_2)_{12}CH_3$, $(CH_2)_{14}CH_3$ and $(CH_2)_7CH=CH(CH_2)_2CH_3$; Y₁ is selected from the group consisting of $(CO)_{1-2}$ and $(CH_2)_{2-7}$;

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and

 Y_2 is selected from the group consisting of CO and CH_2 ;

$$= N^{+}$$

1-myristoyl-2-(4-nitrophenylsuccinyl) phosphatidylcholine (MNP);

$$\rightarrow N^{+}$$
 O P O H S O (d)

$$\begin{array}{c|c} & & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ &$$

wherein

X is selected from the group consisting of O, S, and -O(CO)-; R is selected from the group consisting of (CH₂)₄CH₃, (CH₂)₆CH₃, $(CH_2)_8CH_3$, $(CH_2)_{10}CH_3$, $(CH_2)_{12}CH_3$, $(CH_2)_{14}CH_3$, and (CH₂)₇CH=CH (CH₂)₂CH₃;

 Y_1 is selected from the group consisting of $(CO)_{1-2}$ and $(CH_2)_{2-7}$; and

 Y_2 is selected from the group consisting of CO or CH_2 .

Claim 9 (original): The method of claim 8 where in the substrate is an oxidized derivative of (a), (b), (c), (d) or (e).

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Claim 10-11 (canceled)

Claim 12 (currently amended): The method of claim 1 which further comprising comprises comparing the measured detectable product of step (d) to detectable product in a control comprising an enzymatically active Lp-PLA2 standard.

Claim 13-16 (canceled)

Claim 17 (currently amended): A method for detecting vascular disease in an individual comprising utilizing the method of claim 16 <u>claim 1</u> to determine the individual's Lp-PLA2 activity in a sample wherein increased activity of Lp-PLA2 in the sample is indicative of vascular disease.

Claim 18 (canceled)

Claim 19 (currently amended): A method for selecting an individual for therapy to treat vascular disease comprising utilizing the method of claim 16 claim 1 to determine the individual's Lp-PLA2 activity in a sample wherein increased activity of Lp-PLA2 in the sample is indicative of an individual who will benefit from therapy to treat vascular disease.

Claim 20-21 (canceled)

Claim 22 (currently amended): A method for monitoring an individual's response to therapy to treat vascular

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disease comprising utilizing the method of claim 16 claim 1 to determine the individual's Lp-PLA2 activity in a sample wherein decreased activity of Lp-PLA2 in the sample is indicative of an individual who is responding favorably to therapy to treat vascular disease.

Claim 23-24 (canceled)

Claim 25 (currently amended) A method for measuring enzymatically active Lipoprotein-associated Phospholipase A2 (Lp-PLA2) in a sample comprising:

- (a) contacting an binder a binder, which specifically binds Lp-PLA2, with the sample to form a binder-Lp-PLA2 complex;
- (b) immobilizing the binder-Lp-PLA2 complex;
- (c) washing the immobilized binder-Lp-PLA2 complex to remove an enzymatically active unbound material or an interfering substance(s);
- (d) contacting the immobilized bound Lp-PLA2 with a substrate converted to a detectable product in the presence of Lp-PLA2; and
- (e) measuring detectable product indicative of enzymatically active Lp-PLA2 in the sample.

Claim 26 (original): The method of claim 25, wherein the sample is a serum sample, a plasma sample or an EDTA treated plasma sample.

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Claim 27 (original): The method of claim 25, wherein the binder is an antibody.

Claim 28-29 (canceled)

Claim 30 (currently amended): The method of claim 25 wherein the binder-Lp-PLA2 complex is immobilized by binding to an immobilized compound, said immobilized compound comprising an antibody, protein or compound capable of binding the binder-Lp-PLA2 complex.

Claim 31-34 (canceled)

Claim 35 (original): The method of claim 25 wherein the binder is conjugated to an immobilizing agent.

Claim 36 (original): The method of claim 35, wherein the binder conjugated to an immobilizing agent is an antibody.

Claim 37-38 (canceled)

Claim 39 (original): The method of claim 35 wherein the immobilizing agent is an antibody, protein or compound capable of binding an immobilized compound.

Claim 40-41 (canceled)

Claim 42 (original): The method of claim 35 wherein the immobilizing agent is biotin.

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Claim 43 (original): The method of claim 35 wherein the immobilizing agent, conjugated to the binder-Lp-PLA2 complex, binds to an immobilized compound.

Claim 44 (original) The method of claim 43 wherein the immobilized compound is bound to a multi-well plate, a magnetic bead, or a latex bead.

Claim 45 (original) The method of claim 44 wherein the bound compound is an antibody, protein or compound capable of binding the conjugated immobilizing agent.

Claim 46-47 (canceled)

Claim 48 (original): The method of claim 45 wherein the bound substance is streptavidin.

Claim 49-50 (canceled)

Claim 51 (original): The method of claim 25, wherein the substrate is selected from the group consisting of

wherein,

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X is selected from the group consisting of O, S, and -O(CO)-; R is selected from the group consisting of $(CH_2)_4CH_3$, $(CH_2)_6CH_3$, $(CH_2)_8CH_3$, $(CH_2)_{10}CH_3$, $(CH_2)_{12}CH_3$, $(CH_2)_{14}CH_3$, and $(CH_2)_7CH=CH(CH_2)_2CH_3$;

 Y_1 is selected from the group consisting of $(CO)_{1-2}$ and $(CH_2)_{2-7}$; and

Y2 is selected from the group consisting of CO and CH2;

wherein,

X is selected from the group consisting of O, S, and -O(CO)-; R is selected from the group consisting of $(CH_2)_4CH_3$, $(CH_2)_6CH_3$, $(CH_2)_8CH_3$, $(CH_2)_{10}CH_3$, $(CH_2)_{12}CH_3$, $(CH_2)_{14}CH_3$ and $(CH_2)_7CH=CH(CH_2)_2CH_3$; Y₁ is selected from the group consisting of $(CO)_{1-2}$ and $(CH_2)_{2-7}$; and

 Y_2 is selected from the group consisting of CO and CH_2 ;

1-myristoyl-2-(4-nitrophenylsuccinyl) phosphatidylcholine (MNP);

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$$\rightarrow N^{+}$$
 O $\rightarrow N^{+}$ O $\rightarrow N^$

$$\begin{array}{c|c} & & & & \\ & & & \\ & & & \\ &$$

wherein

X is selected from the group consisting of O, S, and -O(CO)-; R is selected from the group consisting of $(CH_2)_4CH_3$, $(CH_2)_6CH_3$, $(CH_2)_8CH_3$, $(CH_2)_{10}CH_3$, $(CH_2)_{12}CH_3$, $(CH_2)_{14}CH_3$, and $(CH_2)_7CH=CH(CH_2)_2CH_3$;

 Y_1 is selected from the group consisting of (CO) $_{1-2}$ and (CH $_2$) $_{2-7}$; and

 Y_2 is selected from the group consisting of CO or CH_2 .

Claim 52 (original): The method of claim 51 where in the substrate is an oxidized derivative of (a), (b), (c), (d) or (e).

Claim 53-54 (canceled)

Claim 55 (original): The method of claim 25 further comprising comparing the measured detectable product of step (e) to detectable product in a control comprising an enzymatically active Lp-PLA2 standard.

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Claim 56-58 (canceled)

Claim 59 (currently amended): A method for detecting vascular disease in an individual comprising utilizing the method of claim 58 <u>claim 55</u> to determine the individual's Lp-PLA2 activity in a sample wherein increased activity of Lp-PLA2 in the sample is indicative of vascular disease.

Claim 60 (canceled)

Claim 61 (currently amended): A method for selecting an individual for therapy to treat vascular disease comprising utilizing the method of claim 58 claim 55 to determine the individual's Lp-PLA2 activity in a sample wherein increased activity of Lp-PLA2 in the sample is indicative of an individual who will benefit from therapy to treat vascular disease.

Claim 62-63 (canceled)

Claim 64 (currently amended): A method for monitoring an individuals individual's response to therapy to treat vascular disease comprising utilizing the method of claim 58 claim 55 to determine the individual's Lp-PLA2 activity in a sample wherein decreased activity of Lp-PLA2 in the sample is indicative of an individual who is responding favorably to therapy to treat vascular disease.

Claim 65-66 (canceled)

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Claim 67 (original): A kit for measuring enzymatically active Lipoprotein-associated Phospholipase A2 (Lp-PLA2) in a sample comprising a binder which specifically binds Lp-PLA2 and a substrate converted to a detectable product in the presence of Lp-PLA2.

Claim 68 (original): The kit of claim 67 wherein the substrate is selected from the group consisting of

wherein,

X is selected from the group consisting of O, S, and -O(CO)-; R is selected from the group consisting of $(CH_2)_4CH_3$, $(CH_2)_6CH_3$, $(CH_2)_8CH_3$, $(CH_2)_{10}CH_3$, $(CH_2)_{12}CH_3$, $(CH_2)_{14}CH_3$, and $(CH_2)_7CH=CH(CH_2)_2CH_3$;

 Y_1 is selected from the group consisting of $(CO)_{1-2}$ and $(CH_2)_{2-7}$; and

Y2 is selected from the group consisting of CO and CH2;

wherein,

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X is selected from the group consisting of O, S, and -O(CO)-; R is selected from the group consisting of $(CH_2)_4CH_3$, $(CH_2)_6CH_3$, $(CH_2)_8CH_3$, $(CH_2)_{10}CH_3$, $(CH_2)_{12}CH_3$, $(CH_2)_{14}CH_3$ and $(CH_2)_7CH=CH(CH_2)_2CH_3$; Y₁ is selected from the group consisting of $(CO)_{1-2}$ and $(CH_2)_{2-7}$; and

Y2 is selected from the group consisting of CO and CH2;

$$= N^{+}$$

1-myristoyl-2-(4-nitrophenylsuccinyl) phosphatidylcholine (MNP);

$$\begin{array}{c|c} & & & & \\ & & & \\ & & & \\ &$$

wherein

X is selected from the group consisting of O, S, and -O(CO)-; R is selected from the group consisting of $(CH_2)_4CH_3$, $(CH_2)_6CH_3$, $(CH_2)_8CH_3$, $(CH_2)_{10}CH_3$, $(CH_2)_{12}CH_3$, $(CH_2)_{14}CH_3$, and $(CH_2)_7CH=CH(CH_2)_2CH_3$;

 Y_1 is selected from the group consisting of $(CO)_{1-2}$ and $(CH_2)_{2-7}$; and

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 Y_2 is selected from the group consisting of CO or CH_2 .

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Claim 69 (original): The kit of claim 68 wherein the substrate is an oxidized derivative of (a), (b), (c), (d) or (e).

Claim 70 (original): The kit of claim 67 further comprising an enzymatically active Lp-PLA2 standard.

Claim 71-72 (canceled)

Claim 73 (original): A method for measuring enzymatically active Lipoprotein-associated Phospholipase A2 (Lp-PLA2) in a sample comprising:

- (a) incubating the sample with a compound which reduces active thiol(s) in the sample;
- (b) contacting the incubated sample with a substrate converted to a free thiol product in the presence of enzymatically active Lp-PLA2; and
- (c) measuring free thiol product indicative of enzymatically active Lp-PLA2 in the sample.

Claim 74 (original): The method of claim 73, wherein the sample is a serum sample, a plasma sample or an EDTA treated plasma sample.

Claim 75 (canceled)

Claim 76 (currently amended): The method of claim 73 wherein the sample is incubated at room temperature or at 37°C.

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Claim 77 (canceled)

Claim 78 (original): The method of claim 73 wherein the sample is incubated from about 2 to about 120 minutes.

Claim 79 (canceled)

Claim 80 (original): The method of claim 73 wherein the substrate is selected from the group consisting of

2-thio PAF; and

wherein,

R is selected from the group consisting of $(CH_2)_4CH_3$, $(CH_2)_6CH_3$, $(CH_2)_8CH_3$, $(CH_2)_{10}CH_3$, $(CH_2)_{12}CH_3$, $(CH_2)_{14}CH_3$ and $(CH_2)_7CH=CH(CH_2)_2CH_3$; Y_1 is selected from the group consisting of $(CO)_{1-2}$ and $(CH_2)_{2-7}$; and

 Y_2 is selected from the group consisting of CO and CH_2 .

Claim 81 (original): The method of claim 80 where in the

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substrate is an oxidized derivative of (a) or (b).

Claim 82 (original): The method of claim 73 further comprising comparing measured free thiol product of step (c) to free thiol product in a control comprising an enzymatically active Lp-PLA2 standard.

Claim 83-85 (canceled)

Claim 86 (original): A kit for measuring enzymatically active Lipoprotein-associated Phospholipase A2 (Lp-PLA2) in a sample comprising a compound which reduces active thiol(s) and a substrate converted to a detectable product in the presence of Lp-PLA2.

Claim 87 (original): The kit of claim 86 wherein the substrate is selected from the group consisting of

$$\rightarrow N^{+}$$

2-thio PAF; and

$$\begin{array}{c|c} & & & & \\ & & & \\ & & & \\ &$$

wherein,

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R is selected from the group consisting of $(CH_2)_4CH_3$, $(CH_2)_6CH_3$, $(CH_2)_8CH_3$, $(CH_2)_{10}CH_3$, $(CH_2)_{12}CH_3$, $(CH_2)_{14}CH_3$ and $(CH_2)_7CH=CH(CH_2)_2CH_3$; Y_1 is selected from the group consisting of $(CO)_{1-2}$ and $(CH_2)_{2-7}$; and

 Y_2 is selected from the group consisting of CO and CH_2 .

Claim 88 (original): The kit of claim 87 where in the substrate is an oxidized derivative of (a) or (b).

Claim 89 (original): The kit of claim 86 further comprising an enzymatically active Lp-PLA2 standard.

Claim 90-91 (canceled)